

Condensation Reaction of α -Aroyl- α -acetyl Ketene Cyclic Dithioacetals with Aromatic Aldehydes

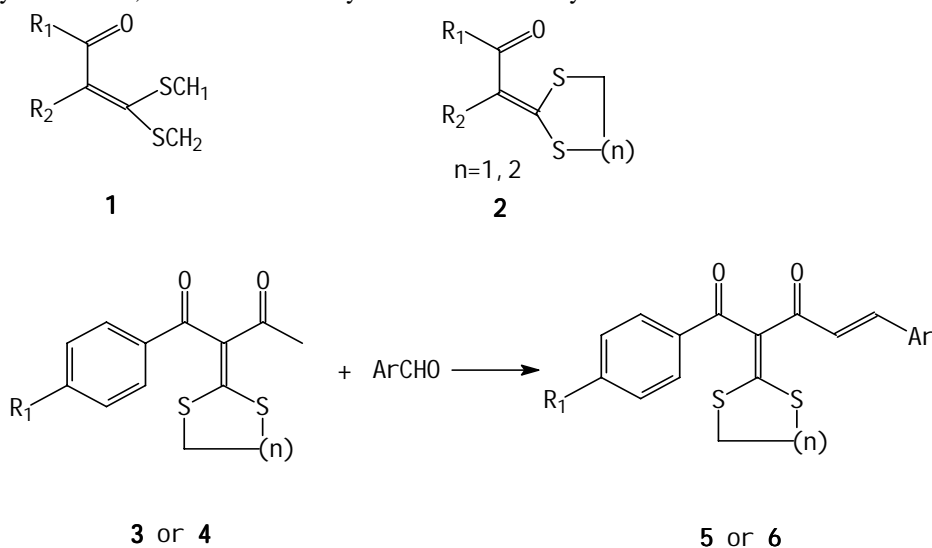
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Abstract: The title compounds **3** and **4** condensed with aromatic aldehydes to give α -aroyl- α -cinnamoyl ketene cyclic dithioacetals **5** and **6** with sodium ethoxide as the base. The stereochemistry of **5** and **6** was assigned as E-configuration by ^1H NMR.

Keywords: α -aroyl- α -cinnamoyl ketene cyclic dithioacetals, aromatic, aldehydes, condensation reaction.

As a versatile three-carbon synthon, α -oxoketene dimethylthio acetals **1** have been applied in many fields^{1,2}. In our previous works³⁻⁵, some properties, especially addition selectivity, of the α -oxoketene cyclic dithioacetals **2** were found to be quite different from those of **1**. Here α -aroyl- α -acetyl ketene cyclic dithioacetals **3** and **4** were allowed to condense with aromatic aldehydes, and fifteen new compounds **5** and **6** were obtained in moderate to high yields. But **5**, **6** from **1** could only be obtained in low yields.



The experiments showed that optimum yields of **5** and **6** were obtained when the temperature of reaction was controlled between 40-60°C.

In our experiments, J between two protons of C=C bond is about 15 Hz and this shows the stereochemistry of **5** and **6** is in E-configuration⁶.

All compounds synthesized are assigned by their IR and ¹H NMR spectra.

Table

Substrate	Product	n	R	Ar	yield (%)
3a	5a	1	H	ph	59
3b	5b	1	H	m-O ₂ N-C ₆ H ₄	52
3c	5c	1	H	p-OCH ₃ C ₆ H ₄	58
3d	5d	1	H		51
3e	5e	1	H	phCH=CH	63
3f	5f	1	H	p-N (CH ₃) ₂ C ₆ H ₄	35
3g	5g	1	CH ₃ O		73
4a	6a	2	H	ph	72
4b	6b	2	H	p-N (CH ₃) ₂ C ₆ H ₄	52
4c	6c	2	H	p-OCH ₃ C ₆ H ₄	80
4d	6d	2	H		74
4e	6e	2	H	phCH=CH	85
4f	6f	2	H		80
4g	6g	2	CH ₃ O	phCH=CH	64
4h	6h	2	CH ₃ O	ph	50

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